

John C Burnett

List of Publications by Year in descending order

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Version: 2024-02-01

41
papers

3,316
citations

257450

24
h-index

361022

35
g-index

45
all docs

45
docs citations

45
times ranked

5167
citing authors

#	ARTICLE	IF	CITATIONS
1	RNA-Based Therapeutics: Current Progress and Future Prospects. <i>Chemistry and Biology</i> , 2012, 19, 60-71.	6.0	804
2	Stochastic Gene Expression in a Lentiviral Positive-Feedback Loop: HIV-1 Tat Fluctuations Drive Phenotypic Diversity. <i>Cell</i> , 2005, 122, 169-182.	28.9	599
3	Current progress of siRNA/shRNA therapeutics in clinical trials. <i>Biotechnology Journal</i> , 2011, 6, 1130-1146.	3.5	380
4	Nanoparticle-Based Delivery of RNAi Therapeutics: Progress and Challenges. <i>Pharmaceuticals</i> , 2013, 6, 85-107.	3.8	171
5	Current Progress of RNA Aptamer-Based Therapeutics. <i>Frontiers in Genetics</i> , 2012, 3, 234.	2.3	111
6	Potent and Targeted Activation of Latent HIV-1 Using the CRISPR/dCas9 Activator Complex. <i>Molecular Therapy</i> , 2016, 24, 488-498.	8.2	109
7	Control of Stochastic Gene Expression by Host Factors at the HIV Promoter. <i>PLoS Pathogens</i> , 2009, 5, e1000260.	4.7	98
8	Combinatorial Latency Reactivation for HIV-1 Subtypes and Variants. <i>Journal of Virology</i> , 2010, 84, 5958-5974.	3.4	97
9	HIV Promoter Integration Site Primarily Modulates Transcriptional Burst Size Rather Than Frequency. <i>PLoS Computational Biology</i> , 2010, 6, e1000952.	3.2	95
10	Dual functional BAFF receptor aptamers inhibit ligand-induced proliferation and deliver siRNAs to NHL cells. <i>Nucleic Acids Research</i> , 2013, 41, 4266-4283.	14.5	73
11	Cell-Specific RNA Aptamer against Human CCR5 Specifically Targets HIV-1 Susceptible Cells and Inhibits HIV-1 Infectivity. <i>Chemistry and Biology</i> , 2015, 22, 379-390.	6.0	71
12	RNA interference approaches for treatment of HIV-1 infection. <i>Genome Medicine</i> , 2015, 7, 50.	8.2	69
13	High throughput sequencing analysis of RNA libraries reveals the influences of initial library and PCR methods on SELEX efficiency. <i>Scientific Reports</i> , 2016, 6, 33697.	3.3	66
14	Molecular basis for improved gene silencing by Dicer substrate interfering RNA compared with other siRNA variants. <i>Nucleic Acids Research</i> , 2013, 41, 6209-6221.	14.5	59
15	Improvements and Limitations of Humanized Mouse Models for HIV Research: NIH/NIAID "Meet the Experts" 2015 Workshop Summary. <i>AIDS Research and Human Retroviruses</i> , 2016, 32, 109-119.	1.1	57
16	AptaTRACE Elucidates RNA Sequence-Structure Motifs from Selection Trends in HT-SELEX Experiments. <i>Cell Systems</i> , 2016, 3, 62-70.	6.2	55
17	The role of antisense long noncoding RNA in small RNA-triggered gene activation. <i>Rna</i> , 2014, 20, 1916-1928.	3.5	46
18	Regulation of host gene expression by HIV-1 TAR microRNAs. <i>Retrovirology</i> , 2013, 10, 86.	2.0	45

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19	HIV Evades RNA Interference Directed at TAR by an Indirect Compensatory Mechanism. <i>Cell Host and Microbe</i> , 2008, 4, 484-494.	11.0	44
20	Exosome-mediated stable epigenetic repression of HIV-1. <i>Nature Communications</i> , 2021, 12, 5541.	12.8	41
21	Creating genetic resistance to HIV. <i>Current Opinion in Immunology</i> , 2012, 24, 625-632.	5.5	39
22	HIV Replication and Latency in a Humanized NSG Mouse Model during Suppressive Oral Combinational Antiretroviral Therapy. <i>Journal of Virology</i> , 2018, 92, .	3.4	36
23	Identification of a therapeutic interfering particle—A single-dose SARS-CoV-2 antiviral intervention with a high barrier to resistance. <i>Cell</i> , 2021, 184, 6022-6036.e18.	28.9	36
24	Aptamer—siRNA Chimeras for HIV. <i>Advances in Experimental Medicine and Biology</i> , 2015, 848, 211-234.	1.6	28
25	Novel Humanized Peripheral Blood Mononuclear Cell Mouse Model with Delayed Onset of Graft-versus-Host Disease for Preclinical HIV Research. <i>Journal of Virology</i> , 2022, 96, JVI0139421.	3.4	11
26	CRISPR-Cas9-mediated gene disruption of HIV-1 co-receptors confers broad resistance to infection in human T cells and humanized mice. <i>Molecular Therapy - Methods and Clinical Development</i> , 2022, 24, 321-331.	4.1	11
27	Stem cells, ribozymes and HIV. <i>Gene Therapy</i> , 2009, 16, 1178-1179.	4.5	10
28	Mutual Information Analysis Reveals Coevolving Residues in Tat That Compensate for Two Distinct Functions in HIV-1 Gene Expression. <i>Journal of Biological Chemistry</i> , 2012, 287, 7945-7955.	3.4	10
29	Arsenic Trioxide and Venetoclax Synergize against AML Progenitors by ROS Induction and Inhibition of Nrf2 Activation. <i>International Journal of Molecular Sciences</i> , 2022, 23, 6568.	4.1	9
30	Progress toward curing HIV infection with hematopoietic cell transplantation. <i>Stem Cells and Cloning: Advances and Applications</i> , 2015, 8, 109.	2.3	8
31	Nucleolar Localization of HIV-1 Rev Is Required, Yet Insufficient for Production of Infectious Viral Particles. <i>AIDS Research and Human Retroviruses</i> , 2018, 34, 961-981.	1.1	7
32	Pre-clinical data supporting immunotherapy for HIV using CMV-HIV-specific CAR T cells with CMV vaccine. <i>Molecular Therapy - Methods and Clinical Development</i> , 2022, 25, 344-359.	4.1	6
33	693. Potent and Targeted Activation of Latent HIV-1 Using Multiplexed Guide RNAs and the CRISPR/dCas9 Activator Complex. <i>Molecular Therapy</i> , 2015, 23, S276.	8.2	5
34	CRED9: a differentially expressed lincRNA regulates expression of transcription factor CEBPA. <i>Rna</i> , 2021, 27, 891-906.	3.5	5
35	Biomolecular Therapeutics for HIV. , 2018, , 541-567.		2
36	Irradiated compared with nonirradiated NSG mice for the development of a human B-cell lymphoma model. <i>Comparative Medicine</i> , 2014, 64, 179-85.	1.0	2

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37	580. In Vivo Analyses of Aptamers and siRNA Therapeutics Against HIV-1 in Humanized Mouse Model. <i>Molecular Therapy</i> , 2015, 23, S231.	8.2	0
38	692. A Novel RNAi Trigger Design Retains Potent, Target Specific Activity Despite Emerging Mutations in the Target Site. <i>Molecular Therapy</i> , 2015, 23, S276.	8.2	0
39	581. Exploring Potency of Small RNAs for HIV Latency Reactivation. <i>Molecular Therapy</i> , 2015, 23, S231.	8.2	0
40	RNA Interference-Based Gene Therapy Strategies for the Treatment of HIV Infection. , 2015, , 1033-1044.		0
41	LGIT In Vitro Latency Model in Primary and T Cell Lines to Test HIV-1 Reactivation Compounds. <i>Methods in Molecular Biology</i> , 2016, 1354, 255-264.	0.9	0